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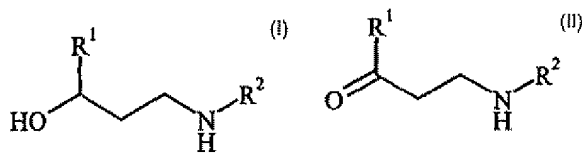
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(54) Title: PROCESS FOR THE PREPARATION OF *N*-MONOSUBSTITUTED β -AMINO ALCOHOLS

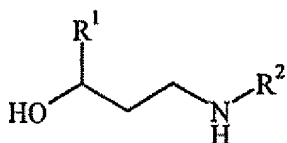


R¹ and R² are as defined above.

(57) Abstract: The invention relates to a process for
the synthesis of *N*-monosubstituted β -amino alcohols
of formula (I) and/or an addition salt of a proton acid,
wherein R¹ and R² independently represent alkyl,
cycloalkyl, aryl or aralkyl, each being optionally further
substituted with alkyl, alkoxy and/or halogen via direct
preparation of *N*-monosubstituted β -amino ketones of
formula and its addition salts of proton acids, wherein

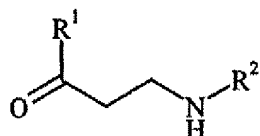
Process for the preparation of *N*-monosubstituted β -amino alcohols

The invention relates to a process for the preparation of *N*-monosubstituted β -amino alcohols of formula



I

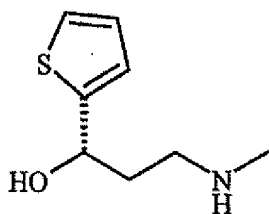
10 and/or an addition salt of a proton acid via direct synthesis of *N*-monosubstituted β -keto amines of formula



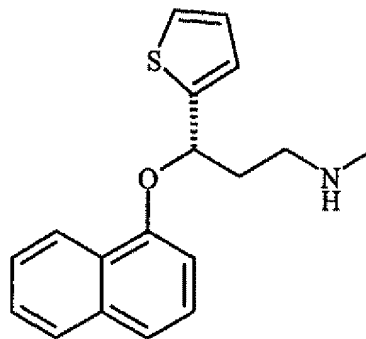
II

and/or an addition salt of a proton acid.

N-Monosubstituted β -amino alcohols of formula I like (*S*)-(-)-3-*N*-methylamino-1-(2-thienyl)-
20 1-propanol (LY293628) are useful key intermediates and building blocks for the preparation of
pharmaceutically active compounds like (*S*)-(+)-methyl-[3-(1-naphthyloxy)-3-(2-thienyl)-
propyl]-amine ((*S*)-duloxetine) (Liu, H. et al., *Chirality* 12 (2000) 26-29), a potential neuro-
active compound which strongly inhibits the serotonin and norephedrine uptake (Deeter, J. et
al., *Tetrahedron Lett.* 31 (1990) 7101-7104).



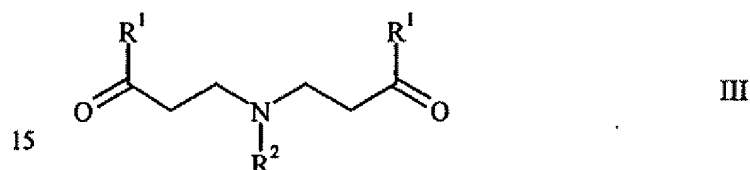
LY293628

(*S*)-duloxetine

In the following the terms "amine" or "amines" include their corresponding addition salts of proton acids.

Direct preparation of *N*-monosubstituted β -keto amines of formula II establishes an alternative
5 and economically advantageous source for industrial production of *N*-monosubstituted β -amino alcohols of formula I.

Compounds of formula II were first synthesized in 1922 by reacting ketones with
formaldehyde and primary or secondary alkylamines in the presence of hydrochloric acid
10 (Mannich, C. et al., *Chem. Ber.* 55 (1922) 356-365). In said reactions with primary alkylamines formation of hydrochlorides of tertiary β -keto amines of formula



prevails over formation of hydrochlorides of secondary β -keto amines of formula II. These findings were supported by Blicke et al. (*J. Am. Chem. Soc.* 64 (1942) 451-454) and Becker et al. (*Wiss. Z. Tech. Hochsch. Chem. Leuna-Merseburg.* 11 (1969) 38-41).

20

According to Mannich et al. steam distillation of tertiary β -keto amines of formula III results in formation of secondary β -keto amines of formula II in fairly satisfactory yields, accompanied by vinyl compounds and other by-products.

25 In spite of the loss of more than 50 % of the starting compounds and due to lack of alternative processes this procedure is still used for the preparation of secondary β -keto amines.

Another drawback in presently known preparation methods of β -keto amines is the need of isolation of the desired intermediate compounds of formula II from unwanted by-products of
30 formula III.

EP-A 457 559 and EP-A 650 965 disclose the preparation of *N,N*-dimethyl β -amino alcohols via Mannich-type reactions of methyl ketones with paraformaldehyde and dimethylamine

followed by reduction of the carbonyl group. After reaction of the hydroxyl group affording alkyl or aryl ether derivatives one methyl radical is removed to obtain *N*-monosubstituted compounds which requires delicate and expensive reactions.

- 5 Only Becker et al. disclose some few examples with yields of about 60% of *N*-monomethyl β -keto amines using *N*-methylammonium oxalates as nitrogen source. Nevertheless, the process disclosed by Becker et al. is not advantageous because it strictly depends on the use of amino oxalates. In contrast to the free amines or corresponding hydrochlorides oxalates of primary amines are not commercially available and their preparation requires further synthesis
10 and purification steps.

Using oxalates is also disadvantageous because it requires additional reduction equivalents in the next step, reducing the ketone intermediates to the title compounds.

- None of the known processes for the production of *N*-monosubstituted β -amino alcohols of
15 formula I and ether derivatives thereof includes, intends or concerns intermediate products comparable to *N*-monosubstituted β -keto amines of formula II of the present invention. Although still many efforts were made to find new preparation processes, the pathway of the present invention for direct synthesis of *N*-monosubstituted β -keto amines and subsequent reduction to *N*-monosubstituted β -amino alcohols is not yet disclosed.

20

- The problem to be solved was to provide an alternative and efficient process for the synthesis of *N*-monosubstituted β -amino alcohols and derivatives thereof in high yields. Furthermore, the proposed process should provide high yields independently of steric aspects of the used amino
25 or carbonyl compounds.

The problems mentioned above could be solved according to claim 1.

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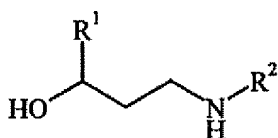
Starting with commercially available methyl ketones and primary amines and/or an addition salt of a proton acid, which were reacted with formaldehyde in the presence a solvent and optionally of a proton acid at a pressure above 1.5 bar *N*-monosubstituted β -amino ketones

which could be directly reduced to the desired *N*-monosubstituted β -amino alcohols were obtained in high yields.

As a further advantage of the instant process high yields of *N*-monomethyl β -amino ketones
 5 can be obtained by direct usage of methylamine hydrochloride which is easily available, cheap and, since it is a solid compound, easy to handle.

The present invention discloses a process for the preparation of a compound of formula

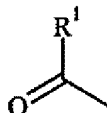
10



I

and/or an addition salt of a proton acid, wherein R^1 and R^2 independently represent alkyl,
 15 cycloalkyl, aryl or aralkyl, each being optionally further substituted with alkyl, alkoxy and/or halogen, which process comprises the steps of
 a) reacting a mixture comprising
 (i) a methyl ketone of formula

20



IV

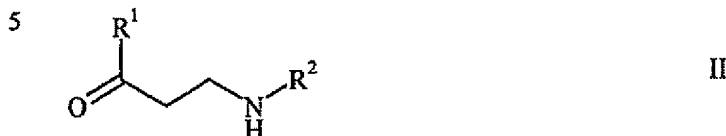
wherein R^1 is as defined above,
 25 (ii) a compound of formula



V

and/or an addition salt of a proton acid, wherein R^2 is as defined above, and
 30 (iii) formaldehyde or a source of formaldehyde selected from the group consisting of formaldehyde in aqueous solution, 1,3,5-trioxane, paraformaldehyde and mixtures thereof, in the presence of
 a solvent selected from the group consisting of water, aliphatic alcohols, cycloaliphatic

alcohols and mixtures thereof, and
optionally a proton acid
to afford a compound of formula



and/or an addition salt of a proton acid, and

10

b) reducing the carbonyl group of said β -amino ketone to afford a compound of formula I,
and/or an addition salt of a proton acid,
wherein the first step is carried out at a pressure above 1.5 bar.

15

In a preferred embodiment R^1 and R^2 can independently represent
linear or branched C_{1-8} alkyl, C_{3-8} cycloalkyl, phenyl, naphthyl, furanyl, benzofuranyl, thienyl,
benzo[b]thienyl or aralkyl, wherein the alkyl moiety of the aralkyl residue is linear C_{1-4} alkyl,
and the aryl moiety is selected from the group consisting of phenyl, naphthyl, furanyl,
benzofuranyl, thienyl and benzo[b]thienyl,

20

each aryl or aralkyl being optionally substituted with halogen, linear or branched C_{1-4} alkyl,
linear or branched C_{1-4} alkoxy, C_{3-6} cycloalkyl, CF_3 , C_2F_5 , OCF_3 or OC_2F_5 .

It is particularly preferred that R^1 represents furanyl or thienyl.

It is also particularly preferred that R^2 represents linear or branched C_{1-8} alkyl. More

25

particularly preferred R^2 represents methyl, ethyl, propyl, isopropyl, butyl, isobutyl or *tert*-
butyl.

Preferably, the compound of formula V is used as a free amine and/or an addition salt of a
proton acid. Particularly preferred are free amines, formates, acetates, oxalates, hydrochlorides,

30

hydrobromides or mixtures thereof. More particularly preferred are free amines and/or
hydrochlorides.

In a preferred embodiment the compound of formula V is present in an amount at least

equimolar to that of the compound of formula IV. Particularly preferred the molar ratio of the compound of formula V to the compound of formula IV is between 1 and 2.

In a preferred embodiment the solvent comprises water, an aliphatic or cycloaliphatic alcohol
5 or a mixture thereof.

Particularly preferred alcohols are linear or branched aliphatic C₁₋₁₂ alcohols, cycloaliphatic C₅₋₈ alcohols, di- and/or trimeric ethylene glycols or mono C₁₋₄ alkyl or acetyl derivatives thereof, each of said alcohols containing 1 to 3 hydroxy groups.

10

Examples for said alcohols are methanol, ethanol, propanol, isopropyl alcohol, butanol, isobutanol, *tert*-butanol, 2-butanol, 1-pentanol, 2-pentanol, 3-pentanol, 1-hexanol, 2-hexanol, cyclopentanol, cyclohexanol, 1,2-ethanediol, 1,2-propanediol, 1,2-butanediol, 2,3-butanediol, 1,4-butanediol, 1,2,3-propanetriol, 1,2,6-hexanetriol, diethylene glycol, diethylene glycol
15 monomethyl ether, diethylene glycol monoethyl ether, diethylene glycol monobutyl ether, diethylene glycol monoacetate, triethylene glycol, triethylene glycol monomethyl ether, triethylene glycol monoethyl ether, triethylene glycol monobutyl ether and triethylene glycol monoacetate.

20 Preferably said alcohol is ethanol, propanol, isopropyl alcohol, butanol, isobutanol, *tert*-butanol, diethylene glycol or triethylene glycol.

The proton acid can be any organic or inorganic acid, the acid being preferably selected from the group consisting of formic acid, acetic acid, propionic acid, oxalic acid, malonic acid,
25 benzoic acid, HF, HCl, HBr, HI, H₂SO₄ and H₃PO₄. In a preferred embodiment the proton acid can be an acidic salt of a polybasic organic or inorganic acid like monoalkali malonates, alkali hydrogensulfates, alkali hydrogenphosphates and alkali hydrogencarbonates.

More preferably the proton acid is selected from the group consisting of formic acid, acetic acid, propionic acid, oxalic acid, HCl and HBr, more preferably it is selected from the group
30 consisting of formic acid, acetic acid, HCl and HBr.

Preferably reaction step a) is carried out either with added addition salts of amines or proton acids, since even distilled free β -amino ketones of formula II tend to decompose and form by-

products while stored, whereas the corresponding addition salts can be stored over a longer period without decomposition. In the products, the ratio of free amine and its salt corresponds to the ratio of added addition salts of amines and proton acids to the whole amine amount during reaction step a).

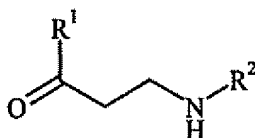
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In a preferred embodiment the pressure during reaction step a) is above 1.5 bar, more preferably in the range of 1.5 to 10 bar and particularly preferred in the range of 1.5 to 5 bar.

In contrast to Becker et al. the inventive process generally allows direct preparation of
 10 *N*-monosubstituted β -keto amines and addition salts of proton acids thereof. The products obtained by the inventive process can be reduced or subsequently reacted without further conversion into other salts.

The present invention also provides a compound of formula

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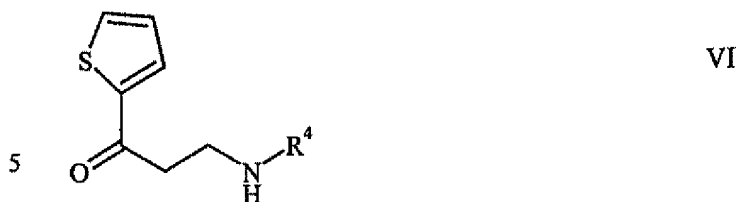
II

and its addition salts of proton acids,

20 wherein R^1 represents furanyl, benzofuranyl, isobenzofuranyl, thienyl or benzo[b]thienyl, each being optionally substituted with halogen, linear or branched C_{1-4} alkyl, linear or branched C_{1-4} alkoxy, C_{3-6} cycloalkyl, CF_3 , C_2F_5 , OCF_3 or OC_2F_5 , and
 wherein R^2 is selected from the group consisting of linear or branched C_{1-8} alkyl, C_{3-8} cycloalkyl, phenyl, naphthyl, furanyl, benzofuranyl, thienyl, benzo[b]thienyl and aralkyl, wherein
 25 the alkyl moiety of the aralkyl residue is linear C_{1-4} alkyl, and the aryl moiety is selected from the group consisting of phenyl, furanyl, benzofuranyl, thienyl and benzo[b]thienyl,
 each aryl or aralkyl being optionally substituted with halogen, linear or branched C_{1-4} alkyl, linear or branched C_{1-4} alkoxy, C_{3-6} cycloalkyl, CF_3 , C_2F_5 , OCF_3 or OC_2F_5 ,
 with the exception of the compound wherein R^1 is thienyl and R^2 is benzyl.

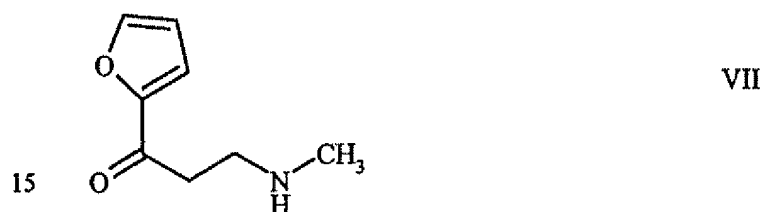
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The present invention also provides a compound of formula



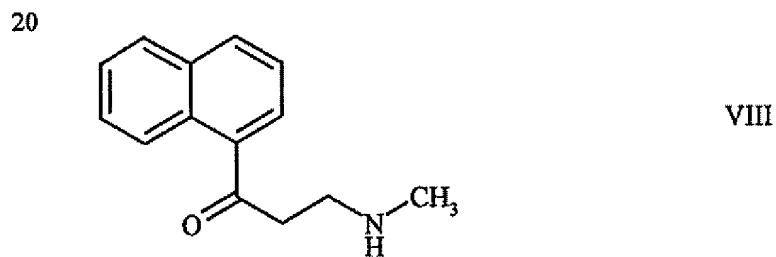
and its addition salts of proton acids, wherein R^4 represents methyl, ethyl, isobutyl and *tert*-butyl.

10 The present invention also provides a compound of formula



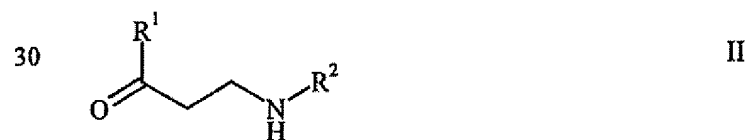
and its addition salts of proton acids.

The present invention also provides a compound of formula



and its addition salts of proton acids.

The present invention also provides a process for the preparation of a compound of formula



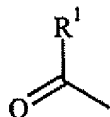
and/or an addition salt of a proton acid, wherein R^1 and R^2 independently represent alkyl,

cycloalkyl, aryl or aralkyl, each being optionally further substituted with alkyl, alkoxy and/or halogen,

which process comprises reacting a mixture comprising

(i) a methyl ketone of formula

5



IV

wherein R¹ is as defined above, and

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(ii) a compound of formula



V

and/or an addition salt of a proton acid, wherein R² is as defined above, and

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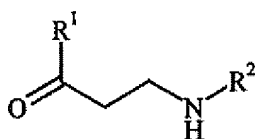
(iii) formaldehyde or a source of formaldehyde selected from the group consisting of formaldehyde in aqueous solution, 1,3,5-trioxane, paraformaldehyde and mixtures thereof, in the presence of

a solvent selected from the group consisting of water, aliphatic alcohols, cycloaliphatic alcohols and mixtures thereof, and

20

optionally a proton acid

to afford a compound of formula



II

25

and/or an addition salt of a proton acid, wherein R¹ and R² are as defined above, and wherein the reaction is carried out at a pressure above 1.5 bar.

In a preferred embodiment R¹ and R² independently represent

30

linear or branched C₁₋₈ alkyl, C₃₋₈ cycloalkyl, phenyl, naphthyl, furanyl, benzofuranyl, thienyl, benzo[b]thienyl and aralkyl, wherein the alkyl moiety of the aralkyl residue is linear C₁₋₄ alkyl, and the aryl moiety is selected from the group consisting of phenyl, naphthyl, furanyl, benzofuranyl, thienyl and benzo[b]thienyl,

each aryl or aralkyl being optionally substituted with halogen, linear or branched C₁₋₄ alkyl, linear or branched C₁₋₄ alkoxy, C₃₋₆ cycloalkyl, CF₃, C₂F₅, OCF₃ or OC₂F₅.

It is particularly preferred that R¹ represents furanyl or thienyl. It is also particularly preferred
5 that R² represents linear or branched C₁₋₈ alkyl. More particularly preferred R² represents methyl, ethyl, propyl, isopropyl, butyl, isobutyl or *tert*-butyl.

Preferably, the compound of formula V can be used as a free amine and/or an addition salt of a
proton acid thereof. Particularly preferred are free amines, formates, acetates, oxalates,
10 hydrochlorides, hydrobromides or mixtures thereof. More particularly preferred are free amines and/or hydrochlorides.

In one preferred embodiment the compound of formula V is present in an amount at least
equimolar to that of the compound of formula IV. Particularly preferred the molar ratio of the
15 compound of formula V to the compound of formula IV is between 1 and 2.

In a preferred embodiment the solvent comprises water, an aliphatic or cycloaliphatic alcohol
or a mixture thereof.

20 Particularly preferred alcohols are linear or branched aliphatic C₁₋₁₂ alcohols, cycloaliphatic C₅₋₈ alcohols, di- and/or trimeric ethylene glycols or mono C₁₋₄ alkyl or acetyl derivatives thereof, each of said alcohols containing 1 to 3 hydroxy groups.

Examples for said alcohols are methanol, ethanol, propanol, isopropyl alcohol, butanol,
25 isobutanol, *tert*-butanol, 2-butanol, 1-pentanol, 2-pentanol, 3-pentanol, 1-hexanol, 2-hexanol, cyclopentanol, cyclohexanol, 1,2-ethanediol, 1,2-propanediol, 1,2-butanediol, 2,3-butanediol, 1,4-butanediol, 1,2,3-propanetriol, 1,2,6-hexanetriol, diethylene glycol, diethylene glycol monomethyl ether, diethylene glycol monoethyl ether, diethylene glycol monobutyl ether, diethylene glycol monoacetate, triethylene glycol, triethylene glycol monomethyl ether,
30 triethylene glycol monoethyl ether, triethylene glycol monobutyl ether and triethylene glycol monoacetate.

Preferably said alcohol is ethanol, propanol, isopropyl alcohol, butanol, isobutanol,

tert-butanol, diethylene glycol or triethylene glycol.

The proton acid can be any organic or inorganic acid, the acid being preferably selected from the group consisting of formic acid, acetic acid, propionic acid, oxalic acid, malonic acid, benzoic acid, HF, HCl, HBr, HI, H₂SO₄ and H₃PO₄. In a preferred embodiment the proton acid is an acidic salt of a polybasic organic or inorganic acids like monoalkali malonates, alkali hydrogensulfates, alkali hydrogenphosphates and alkali hydrogencarbonates. More preferably the proton acid is selected from the group consisting of formic acid, acetic acid, propionic acid, oxalic acid, HCl and HBr, more preferably it is selected from the group consisting of formic acid, acetic acid, HCl and HBr.

In a preferred embodiment the pressure during the reaction is above 1.5 bar, more preferably in the range of 1.5 to 10 bar and particularly preferred in the range of 1.5 to 5 bar.

The present invention is illustrated by the following non-limiting examples.

General Procedure for Examples 1 to 8

A mixture of methyl ketone (1 equivalent (eq)), primary alkyl amine and/or an addition salt thereof (1.1 to 1.5 eq), formaldehyde (1.4 to 1.5 eq), a solvent, optionally in the presence of a proton acid, is heated in an autoclave at a total pressure above 1.5 bar for 5 to 24 hours. Afterwards, the reaction solution is cooled to 20 °C. Optionally the reaction solvent can then be removed partly or in whole and a solvent like ethyl acetate or isopropyl alcohol can be added under vigorous stirring, if necessary to facilitate precipitation of the product. The suspension is cooled (0 to 20 °C) and filtered after precipitation (0.5 to 10 hours), optionally washed and dried to afford a slightly yellow to white powder in a yield between 50 and 75 %. The product can be recrystallized from isopropyl alcohol and/or ethyl acetate if necessary. If the stability of the free base is sufficient at ambient conditions, extracting with an organic solvent and an aqueous base affords the free base.

General Procedure for Comparative Examples 1 to 6

A mixture of methyl ketone (1 eq), primary alkyl amine and/or an addition salt thereof (1 to 1.5 eq), formaldehyde (1.0 to 1.5 eq), optionally in the presence of a proton acid, is heated in refluxing solvent for 5 to 24 hours. Afterwards, the mixture is cooled to 20 °C. Optionally

the reaction solvent can then be removed partly or in whole and a solvent like ethyl acetate or isopropyl alcohol can be added under vigorous stirring, if necessary to facilitate precipitation of the product. The suspension is cooled (0 to 20 °C) and filtered after precipitation (0.5 to 10 hours), optionally washed and dried to afford a slightly yellow to white powder in a yield
5 between 30 and 45 %. The product can be recrystallized from isopropyl alcohol and/or ethyl acetate if necessary.

Example 1: 3-(Methylamino)-1-(thiophen-2-yl)propan-1-one hydrochloride (II, R¹ = thiophen-2-yl, R² = methyl)

10 2-Acetylthiophene (25.5 g, 200 mmol); methylamine hydrochloride (14.9 g, 220 mmol, 1.1 eq); paraformaldehyde (8.2 g, 280 mmol, 1.4 eq); HCl conc. (1.0 g); ethanol (100 mL); 110 °C for 9 hours; ca. 2 to 2.5 bar; removing of ethanol (50 mL) *in vacuo*; addition of ethyl acetate (200 mL); ca. 71 % yield.

¹H-NMR δ (DMSO-d₆, 400 MHz): 9.16 (2 H, s, br), 8.07 (1 H, dd, *J* = 5.0, 1.0), 8.01 (1 H, dd, *J* = 3.8, 1.0), 7.29 (1 H, dd, *J* = 5.0, 3.8), 3.49 (2 H, t), 3.20 (2 H, t), 2.56 (3 H, s).

15 ¹³C-NMR δ (DMSO-d₆, 100 MHz): 189.9, 142.7, 135.4, 133.8, 128.8, 43.1, 34.6, 32.4.

Example 2: 3-(Methylamino)-1-(thiophen-2-yl)propan-1-one hydrochloride (II, R¹ = thiophen-2-yl, R² = methyl)

20 2-Acetylthiophene (24.9 g, 197 mmol); methylamine hydrochloride (14.8 g, 219 mmol, 1.1 eq); paraformaldehyde (8.3 g, 276 mmol, 1.4 eq); HCl conc. (1.1 g); isopropyl alcohol (100 mL); 110 °C for 8 hours; ca. 2 to 2.5 bar; addition of isopropyl alcohol (50 mL); ca. 65 % yield.

25 **Comparative Example 1:** 3-(Methylamino)-1-(thiophen-2-yl)propan-1-one hydrochloride (II, R¹ = thiophen-2-yl, R² = methyl)

2-Acetylthiophene (7.9 g, 300 mmol); methylamine hydrochloride (30.4 g, 450 mmol, 1.5 eq); paraformaldehyde (12.6 g, 420 mmol, 1.4 eq); HCl conc. (1.5 g); isopropyl alcohol (200 mL); heating under reflux (82 °C) for 8 hours; addition of ethyl acetate (200 mL); ca. 43 % yield.

30

Example 3: 3-(Ethylamino)-1-(thiophen-2-yl)propan-1-one hydrochloride (II, R¹ = thiophen-2-yl, R² = ethyl)

2-Acetylthiophene (6.3 g, 50 mmol); ethylamine hydrochloride (6.1 g, 75 mmol, 1.5 eq);

paraformaldehyde (2.1 g, 75 mmol, 1.5 eq); HCl conc. (0.3 g); ethanol (35 mL); 110 °C for 9 hours; ca. 2 to 2.5 bar; removing of ethanol (25 mL) *in vacuo*; addition of ethyl acetate (50 mL); ca. 73 % yield.

¹H-NMR δ (DMSO-d₆, 400 MHz): 9.3 (2 H, s, br), 8.08 (1 H, dd), 8.00 (1 H, dd), 7.28

5 (1 H, dd), 3.51 (2 H, t), 3.20 (2 H, t), 2.96 (2 H, q), 1.23 (3 H, t).

Comparative Example 2: 3-(Ethylamino)-1-(thiophen-2-yl)propan-1-one hydrochloride (II, R¹ = thiophen-2-yl, R² = ethyl)

2-Acetylthiophene (12.6 g, 100 mmol); ethylamine hydrochloride (12.2 g, 150 mmol, 1.5 eq);
10 paraformaldehyde (4.1 g, 140 mmol, 1.4 eq); HCl conc. (0.5 g); ethanol (70 mL); heating under reflux (78 °C) for 6 hours; removing of ethanol (25 mL) *in vacuo*; addition of ethyl acetate (70 mL); ca. 31 % yield.

Example 4: 3-(Isobutylamino)-1-(thiophen-2-yl)propan-1-one hydrochloride (II, R¹

15 = thiophen-2-yl, R² = isobutyl)

2-Acetylthiophene (6.3 g, 50 mmol); isobutylamine hydrochloride (8.3 g, 75 mmol, 1.5 eq); paraformaldehyde (2.1 g, 75 mmol, 1.5 eq); HCl conc. (0.3 g); ethanol (35 mL); 110 °C for 9 hours; ca. 2 to 2.5 bar; removing of ethanol (35 mL) *in vacuo*; addition of ethyl acetate (50 mL); ca. 56 % yield.

20 ¹H-NMR δ (DMSO-d₆, 400 MHz): 9.0 (2 H, s, br), 8.08 (1 H, dd), 7.99 (1 H, dd), 7.29 (1 H, dd), 3.55 (2 H, t), 3.22 (2 H, t), 2.78 (2 H, d), 2.03 (1 H, m), 0.96 (6 H, d).

Comparative Example 3: 3-(Isobutylamino)-1-(thiophen-2-yl)propan-1-one hydrochloride (II, R¹ = thiophen-2-yl, R² = isobutyl)

25 2-Acetylthiophene (12.6 g, 100 mmol); isobutylamine hydrochloride (16.5 g, 150 mmol, 1.5 eq); paraformaldehyde (4.1 g, 140 mmol, 1.4 eq); HCl conc. (0.5 g); butanol (70 mL); heating under reflux (108 °C) for 7 hours; addition of ethyl acetate (100 mL); ca. 40 % yield.

Example 5: 3-(*tert*-Butylamino)-1-(thiophen-2-yl)propan-1-one hydrochloride (II, R¹ =

30 thiophen-2-yl, R² = *tert*-butyl)

2-Acetylthiophene (6.3 g, 50 mmol); *tert*-butylamine hydrochloride (8.3 g, 75 mmol, 1.5 eq); paraformaldehyde (2.1 g, 75 mmol, 1.5 eq); HCl conc. (0.3 g); butanol (35 mL); 117 °C for 9 hours; ca. 2 to 2.5 bar; addition of ethyl acetate (50 mL); ca. 52 % yield.

¹H-NMR δ (DMSO-d₆, 400 MHz): 9.2 (2 H, s, br), 8.08 (1 H, dd), 7.98 (1 H, dd), 7.30 (1 H, dd), 3.54 (2 H, t), 3.19 (2 H, t), 1.34 (9 H, s).

Comparative Example 4: 3-(*tert*-Butylamino)-1-(thiophen-2-yl)propan-1-one hydrochloride

5 (II, R¹ = thiophen-2-yl, R² = *tert*-butyl)

2-Acetylthiophene (12.6 g, 100 mmol); *tert*-butylamine hydrochloride (16.5 g, 150 mmol, 1.5 eq); paraformaldehyde (4.1 g, 140 mmol, 1.4 eq); HCl conc. (0.5 g); butanol (70 mL); heating under reflux (108 °C) for 18 hours; addition of ethyl acetate (100 mL); ca. 37 % yield.

10 **Example 6:** 3-(Methylamino)-1-(furan-2-yl)propan-1-one hydrochloride (II, R¹ = furan-2-yl, R² = methyl)

2-Acetylfuran (7.5 g, 68 mmol); methylamine hydrochloride (6.9 g, 102 mmol, 1.5 eq); paraformaldehyde (3.1 g, 102 mmol, 1.5 eq); HCl conc. (1.15 g); ethanol (35 mL); 110 °C for 8 hours; ca. 2 to 2.5 bar; removing of ethanol (30 mL) *in vacuo*; addition of ethyl acetate

15 (50 mL); ca. 64 % yield.

¹H-NMR δ (DMSO-d₆, 400 MHz): 9.0 (2 H, s, br), 8.05 (1 H, m), 7.53 (1 H, m), 6.77 (1 H, m), 3.34 (2 H, t), 3.2 (2 H, m), 2.57 (3 H, s, br).

Comparative Example 5: 3-(Methylamino)-1-(furan-2-yl)propan-1-one hydrochloride (II,

20 R¹ = furan-2-yl, R² = methyl)

2-Acetylfuran (11.0 g, 100 mmol); methylamine hydrochloride (10.1 g, 150 mmol, 1.5 eq); paraformaldehyde (4.1 g, 140 mmol, 1.4 eq); HCl conc. (0.5 g); butanol (70 mL); heating under reflux (108 °C) for 7 hours; addition of ethyl acetate (100 mL); ca. 44 % yield.

25 **Example 7:** 3-(Methylamino)-1-phenylpropan-1-one hydrochloride (II, R¹ = phenyl, R² = methyl)

2-Acetophenone (21.0 g, 175 mmol); methylamine hydrochloride (17.5 g, 263 mmol, 1.5 eq); paraformaldehyde (7.9 g, 263 mmol, 1.5 eq); HCl conc. (1.1 g); ethanol (130 mL); 115 °C for 24 hours; ca. 2 to 2.5 bar; addition of ethyl acetate (170 mL); ca. 52 % yield.

30 ¹H-NMR δ (DMSO-d₆, 400 MHz): 9.2 (2 H, s, br), 8.0 (2 H, m), 7.7 (1 H, m), 7.6 (2 H, m), 3.55 (2 H, t), 3.21 (2 H, t), 2.59 (3 H, s).

Example 8: 3-(Methylamino)-1-(2-naphthyl)propan-1-one hydrochloride (II, R¹ = 2-naphthyl,

R² = methyl)

2-Acetonaphthone (8.5 g, 50 mmol); methylamine hydrochloride (5.1 g, 75 mmol, 1.5 eq); paraformaldehyde (2.1 g, 75 mmol, 1.5 eq); HCl conc. (0.3 g); ethanol (35 mL); 117 °C for 14 hours; ca. 2 to 2.5 bar; removing of ethanol (35 mL) *in vacuo*; addition of ethyl acetate

5 (50 mL); ca. 60 % yield.

¹H-NMR δ (DMSO-d₆, 400 MHz): 9.3 (2 H, s, br), 8.74 (1 H, s), 8.17 (1 H, d), 8.0 (3 H, m), 7.7 (2 H, m), 3.70 (2 H, t), 3.28 (2 H, m), 2.60 (3 H, s).

Comparative Example 6: 3-(Methylamino)-1-(2-naphthyl)propan-1-one hydrochloride (II, R¹
10 = 2-naphthyl, R² = methyl)

2-Acetonaphthone (17.0 g, 100 mmol); methylamine hydrochloride (10.1 g, 150 mmol, 1.5 eq); paraformaldehyde (4.1 g, 140 mmol, 1.4 eq); HCl conc. (0.5 g); ethanol (70 mL); heating under reflux (78 °C) for 5 hours; removing of ethanol (30 mL) *in vacuo*; addition of ethyl acetate (100 mL); ca. 42 % yield.

15

Example 9: 3-(Methylamino)-1-(thiophen-2-yl)propan-1-ol (I, R¹ = thiophen-2-yl, R² = methyl)

To a mixture of 3-(methylamino)-1-(thiophen-2-yl)propan-1-one hydrochloride (10.3 g, 50 mmol) and ethanol (35 mL) at 4 °C sodium hydroxide (4.0 g of a 50 % aqueous solution)
20 was added in about 5 minutes. Afterwards, neat sodium borohydride (0.95 g, 25 mmol, 1.0 eq) was added in several portions in about 30 minutes. At the end of the addition, the suspension was stirred for 4 h at the same temperature, then acetone (10.0 mL) was added dropwise in 5 minutes and the mixture was stirred for 10 additional minutes. Water (20 mL) was then added. Afterwards, the mixture was concentrated about 5 times under vacuum and the residue was
25 extracted with tert-butyl methyl ether (2 x 20 mL). The collected organic phases were finally concentrated under vacuum affording an orange oil which crystallised spontaneously after a few hours. Finally, an orange solid was obtained (7.2 g, 84 % yield). This compound can then be used without further purification.

¹H-NMR δ (DMSO-d₆, 400 MHz): 7.35 (1 H, dd, J = 4.8, 1.0), 6.94 (1 H, dd, J = 4.8, 3.6),
30 6.90 (1 H, dd, J = 3.6, 1.0), 4.90 (1 H, t), 3.7 (2 H, m), 2.56 (2 H, m), 2.25 (3 H, s), 1.79 (2 H, q).

¹³C-NMR δ (DMSO-d₆, 100 MHz): 150.9, 126.3, 123.7, 122.3, 67.8, 48.5, 38.7, 36.0.

Example 10: 3-(Isobutylamino)-1-(thiophen-2-yl)propan-1-ol (I, R¹ = thiophen-2-yl, R² = methyl)

To a mixture of 3-(isobutylamino)-1-(thiophen-2-yl)propan-1-one hydrochloride (4.2 g, 19.4 mmol) and ethanol (10 mL) at 4 °C sodium hydroxide (1.6 g of a 50 % aqueous solution) was added in about 20 minutes. Afterwards, neat sodium borohydride (0.37 g, 9.7 mmol, 1.0 eq) was added in several portions in about 30 minutes. At the end of the addition, the suspension was stirred for 4 h at the same temperature, then acetone (10.0 mL) was added dropwise in 20 minutes and the mixture was stirred for 10 additional minutes. Afterwards the precipitate was removed by filtration and the mixture was concentrated under vacuum affording an orange oil.

10 The crude product was purified by column chromatography using a 40 : 10 : 1 (v : v : v) mixture of methylene chloride/methanol/ammonium hydroxide (25 % aqueous solution) affording 3.1 g (76 % yield) of product.

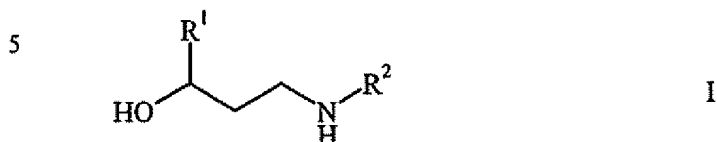
¹H-NMR δ (DMSO-d₆, 400 MHz): 7.20 (1 H, dd, J = 4.8, 1.0), 6.98 (1 H, dd), 6.94 (1 H, dd, J = 4.8, 3.6), 5.20 (1 H, dd), 4.98 (2 H, br), 3.02 (1 H, m), 2.93 (1 H, m), 2.43 (2H, symm. m), 2.03 (1 H, m), 1.97 (1 H, m), 1.80 (1 H, sept), 0.95 (6 H, d).

15

¹³C-NMR δ (DMSO-d₆, 100 MHz): 150.9, 126.3, 123.8, 122.5, 72.1, 57.8, 48.5, 37.4, 28.2, 20.8.

Claims

1. A process for the preparation of a compound of formula



- 10 and/or an addition salt of a proton acid, wherein R¹ and R² independently represent alkyl, cycloalkyl, aryl or aralkyl, each aryl or aralkyl being optionally further substituted with alkyl, alkoxy and/or halogen, which process comprises the following steps

a) reacting a mixture comprising

- (i) a methyl ketone of formula

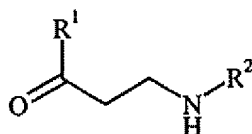
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- 20 wherein R¹ is as defined above, and
(ii) a compound of formula



- 25 and/or an addition salt of proton acid, wherein R² is as defined above, and
(iii) formaldehyde or a source of formaldehyde selected from the group consisting of formaldehyde in aqueous solution, 1,3,5-trioxane, paraformaldehyde and mixtures thereof, in the presence of
a solvent selected from the group consisting of water, aliphatic alcohols,
30 cycloaliphatic alcohols and mixtures thereof, and
optionally a proton acid
to afford a β-amino ketone of formula



II

and/or an addition salt of a proton acid, and

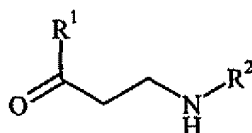
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b) reducing the carbonyl group of said β -amino ketone to afford a compound of formula I, and/or an addition salt of a proton acid

wherein the first step is carried out at a pressure above 1.5 bar.

- 10 2. The process of claim 1 wherein R^1 is selected from the group consisting of linear or branched C_{1-8} alkyl, C_{3-8} cycloalkyl, phenyl, naphthyl, furanyl, benzofuranyl, thienyl, benzo[b]thienyl and aralkyl, wherein the alkyl moiety of the aralkyl residue is linear C_{1-4} alkyl, and the aryl moiety is selected from the group consisting of phenyl, naphthyl, furanyl, benzofuranyl, thienyl and benzo[b]thienyl,
 - 15 each aryl or aralkyl being optionally substituted with halogen, linear or branched C_{1-4} alkyl, linear or branched C_{1-4} alkoxy, C_{3-6} cycloalkyl, CF_3 , C_2F_5 , OCF_3 or OC_2F_5 .
- 20 3. The process of claim 1 or 2 wherein R^2 is selected from the group consisting of linear or branched C_{1-8} alkyl, C_{3-8} cycloalkyl, phenyl, naphthyl, furanyl, benzofuranyl, thienyl, benzo[b]thienyl and aralkyl, wherein the alkyl moiety of the aralkyl residue is linear C_{1-4} alkyl, and the aryl moiety is selected from the group consisting of phenyl, naphthyl, furanyl, benzofuranyl, thienyl and benzo[b]thienyl,
 - 25 each aryl or aralkyl being optionally substituted with halogen, linear or branched C_{1-4} alkyl, linear or branched C_{1-4} alkoxy, C_{3-6} cycloalkyl, CF_3 , C_2F_5 , OCF_3 or OC_2F_5 .
4. The process of any of claims 1 to 3, wherein the compound of formula V is present in an amount at least equimolar to that of the compound of formula IV.
- 30 5. The process of any of claims 1 to 4, wherein the proton acid is a carboxylic or an inorganic acid, the acid being preferably selected from the group consisting of formic acid, acetic acid, propionic acid, oxalic acid, malonic acid, benzoic acid, HF, HCl, HBr, HI, H_2SO_4 , H_3PO_4 , mono alkali malonate, alkali hydrogensulfates, alkali hydrogenphosphates and alkali hydrogencarbonates.

6. The process of any of claims 1 to 5, wherein aliphatic and cycloaliphatic alcohols are selected from the group selected of linear or branched aliphatic C₁₋₁₂ alcohols, cycloaliphatic C₅₋₈ alcohols, di- and/or triethylene glycols and mono C₁₋₄ alkyl or acetyl derivatives thereof, each of said alcohols containing 1 to 3 hydroxy groups.
7. The process of claim 6, wherein the alcohol is selected from the group consisting of methanol, ethanol, propanol, isopropyl alcohol, butanol, isobutanol, *tert*-butanol, 1-pentanol, 2-pentanol, 3-pentanol, 1-hexanol, 2-hexanol, cyclopentanol, cyclohexanol, 1,2-ethanediol, 1,2-propanediol, 1,2-butanediol, 2,3-butanediol, 1,4-butanediol, 1,2,3-propanetriol, 1,2,6-hexanetriol, diethylene glycol, diethylene glycol monomethyl ether, diethylene glycol monoethyl ether, diethylene glycol monobutyl ether, diethylene glycol monoacetate, triethylene glycol, triethylene glycol monomethyl ether, triethylene glycol monoethyl ether, triethylene glycol monobutyl ether and triethylene glycol monoacetate.
8. The process of any of claims 1 to 7, wherein the pressure during reaction step a) is above 1.5 bar, more preferably in the range of 1.5 to 10 bar and more particularly preferred in the range of 1.5 to 5 bar.
9. A compound of formula

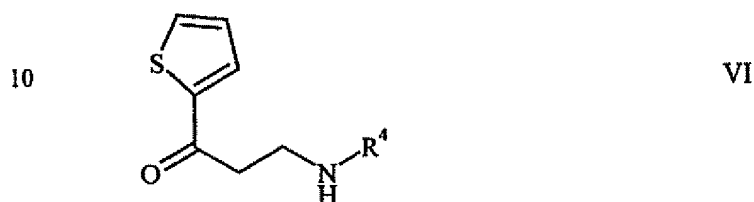


II

- and its addition salts of proton acids, wherein R¹ represents furanyl, benzofuranyl, isobenzofuranyl, thienyl or benzo[b]thienyl, each being optionally substituted with halogen, linear or branched C₁₋₄ alkyl, linear or branched C₁₋₄ alkoxy, C₃₋₆ cycloalkyl, CF₃, C₂F₅, OCF₃ or OC₂F₅; and wherein R² is selected from the group consisting of linear or branched C₁₋₈ alkyl, C₃₋₈ cycloalkyl, phenyl, naphthyl, furanyl, benzofuranyl, thienyl, benzo[b]thienyl and aralkyl, wherein the alkyl moiety of the aralkyl residue is linear C₁₋₄ alkyl, and the aryl moiety is selected from the group consisting of phenyl,

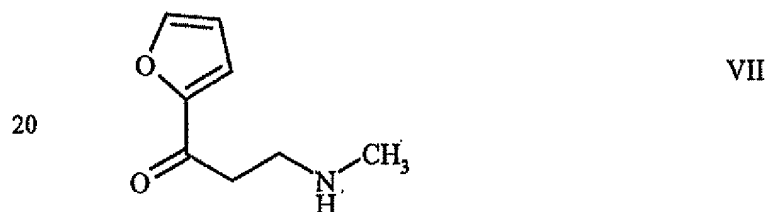
naphthyl, furanyl, benzofuranyl, thienyl and benzo[b]thienyl,
each aryl or aralkyl being optionally substituted with halogen, linear or branched
C₁₋₄ alkyl, linear or branched C₁₋₄ alkoxy, C₃₋₆ cycloalkyl, CF₃, C₂F₅, OCF₃ or OC₂F₅
with the exception of the compound wherein R¹ represents thienyl and R² represents
5 benzyl.

10. A compound of formula



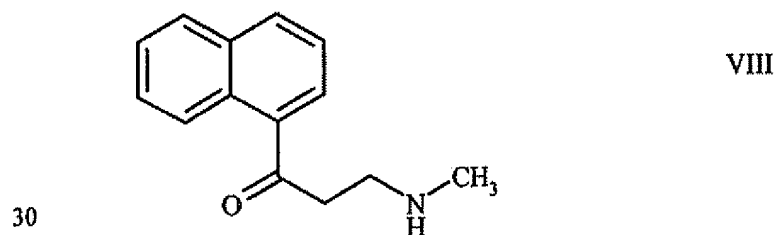
and its addition salts of proton acids, wherein R⁴ represents methyl, ethyl, isobutyl or *tert*-
15 butyl.

11. A compound of formula



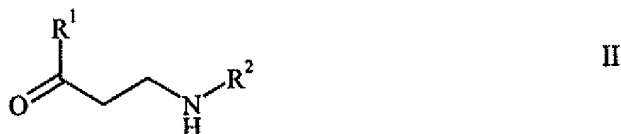
and its addition salts of proton acids.

25 12. A compound of formula



and its addition salts of proton acids.

13. A process for the preparation of a compound of formula



5

and/or an addition salt of a proton acid, wherein R^1 and R^2 independently represent alkyl, cycloalkyl, aryl or aralkyl, each being optionally further substituted with alkyl, alkoxy and/or halogen, which process comprises reacting

(i) a methyl ketone of formula

10



15

wherein R^1 is as defined above, and

(ii) a compound of formula



20

and/or an addition salt of a proton acid, wherein R^2 is as defined above, and

(iii) formaldehyde or a source of formaldehyde selected from the group consisting of formaldehyde in aqueous solution, 1,3,5-trioxane, paraformaldehyde and mixtures thereof, in the presence of

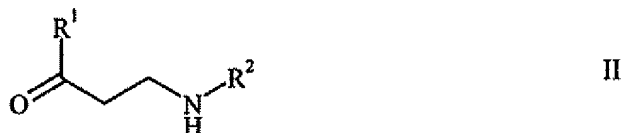
a solvent selected from the group consisting of water, aliphatic alcohols, cycloaliphatic alcohols and mixtures thereof, and

25

optionally a proton acid

to afford a β -amino ketone of formula

30



and/or an addition salt of a proton acid, wherein R^1 and R^2 are as defined above, and

wherein the reaction is carried out at a pressure above 1.5 bar.

14. The process of claim 13 wherein R^1 is as defined in claim 2.
- 5 15. The process of claim 13 or 14 wherein R^2 is as defined in claim 3.
16. The process of any of claims 13 to 15, wherein the compound of formula V is present in an amount at least equimolar to that of the compound of formula IV.
- 10 17. The process of any of claims 13 to 16, wherein the proton acid is a carboxylic or an inorganic acid, preferably the acid is selected from the group consisting of formic acid, acetic acid, propionic acid, oxalic acid, malonic acid, benzoic acid, HF, HCl, HBr, HI, H_2SO_4 , H_3PO_4 , mono alkali malonate, alkali hydrogensulfates, alkali hydrogenphosphates and alkali hydrogencarbonates.
- 15 18. The process of any of claims 16 to 17, wherein aliphatic and cycloaliphatic alcohols are selected from the group consisting of linear or branched aliphatic C_{1-12} alcohols, cycloaliphatic C_{5-8} alcohols, di- triethylene glycols and mono C_{1-4} alkyl or acetyl derivatives thereof, each of said alcohols containing 1 to 3 hydroxy groups.
- 20 19. The process of claim 18, wherein the alcohol is selected from the group consisting of methanol, ethanol, propanol, isopropyl alcohol, butanol, isobutanol, *tert*-butanol, 1-pentanol, 2-pentanol, 3-pentanol, 1-hexanol, 2-hexanol, cyclopentanol, cyclohexanol, 1,2-ethanediol, 1,2-propanediol, 1,2-butanediol, 2,3-butanediol, 1,4-butanediol, 25 1,2,3-propanetriol, 1,2,6-hexanetriol, diethylene glycol; diethylene glycol monomethyl ether, diethylene glycol monoethyl ether, diethylene glycol monobutyl ether, diethylene glycol monoacetate, triethylene glycol, triethylene glycol monomethyl ether, triethylene glycol monoethyl ether, triethylene glycol monobutyl ether and triethylene glycol monoacetate.
- 30 20. The process of any of claims 13 to 19, wherein the pressure during the reaction is above 1.5 bar, more preferably in the range of 1.5 to 10 bar and more particularly preferred in the range of 1.5 to 5 bar.

INTERNATIONAL SEARCH REPORT

International Application No.

PCT/EP 03/07411

A. CLASSIFICATION OF SUBJECT MATTER		
IPC 7 C07C213/08 C07D307/46 C07D333/22 C07C225/12 C07C221/00		
According to International Patent Classification (IPC) or to both national classification and IPC		
B. FIELDS SEARCHED		
Minimum documentation searched (classification system followed by classification symbols)		
IPC 7 C07C C07D		
Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched		
Electronic data base consulted during the international search (name of data base and, where practical, search terms used)		
EPO-Internal, CHEM ABS Data		
C. DOCUMENTS CONSIDERED TO BE RELEVANT		
Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
A	EP 0 457 559 A (LILLY CO ELI) 21 November 1991 (1991-11-21) cited in the application the whole document ---	1-9,13
A	EP 0 650 965 A (LILLY CO ELI) 3 May 1995 (1995-05-03) cited in the application the whole document ---	1-9,13
A	EP 0 046 288 A (RUHRCHEMIE AG) 24 February 1982 (1982-02-24) page 9, line 18 - line 27; example 1 ---	1
A	F. F. BLICKE: "The Mannich reaction" ORGANIC REACTIONS, vol. I, 1942, pages 303-341, XP002218609 the whole document ---	1
-/-		
<input checked="" type="checkbox"/> Further documents are listed in the continuation of box C. <input checked="" type="checkbox"/> Patent family members are listed in annex.		
* Special categories of cited documents: "A" document defining the general state of the art which is not considered to be of particular relevance "E" earlier document but published on or after the international filing date "L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified) "O" document referring to an oral disclosure, use, exhibition or other means "P" document published prior to the international filing date but later than the priority date claimed "T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention "X" document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone "Y" document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art "8" document member of the same patent family		
Date of the actual completion of the international search		Date of mailing of the international search report
27 October 2003		03/11/2003
Name and mailing address of the ISA European Patent Office, P.B. 5818 Patentlaan 2 NL - 2280 HV Rijswijk Tel. (+31-70) 340-2040, Tx. 31 651 epo nl, Fax: (+31-70) 340-3016		Authorized officer Rufet, J

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PCT/EP 03/07411

C.(Continuation) DOCUMENTS CONSIDERED TO BE RELEVANT		
Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
A	<p>KIYOSHI MATSUMOTO: "Synthese unter hohem Druck: Mannich-Reaktion....." ANGEWANDTE CHEMIE., vol. 94, no. 12, 1982, page 937 XP002218608 VCH VERLAGSGESELLSCHAFT, WEINHEIM., DE ISSN: 0044-8249 page 937</p>	1
X	<p>CHEMICAL ABSTRACTS, vol. 56, no. 1, 1962 Columbus, Ohio, US; abstract no. 363g, G. I. DENIS ET AL.: "Alkylation of aromatic amines by the Mannich bases." page 363; column 1; XP002218610 abstract & IZVEST. VYSSHIKH UCHEB. ZAVEDENII, KHIM. I KHIM. TEKHNOL., vol. 4, 1961, pages 426-428,</p>	9
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X	<p>CHEMICAL ABSTRACTS, vol. 63, no. 8, 1965 Columbus, Ohio, US; abstract no. 9900g, R. LANDI-VITTORY ET AL.: "Derivatives of benzofuran and coumaran." page 9899; column 2; XP002218612 abstract & FARMACO (PAVIA), vol. 18, no. 2, 1965, pages 109-118,</p> <p style="text-align: center;">-/--</p>	9

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X	CHEMICAL ABSTRACTS, vol. 67, no. 11, 1967 Columbus, Ohio, US; abstract no. 53946c, B. I. ARDASHEV ET AL.: "Synthesis of Beta-arylamino ketones of the furan series." page 5066; column 1; XP002218613 abstract & KHIM. GETEROTSIKL. SOEDIN., vol. 1, 1967, pages 7-9, ---	9
X	CHEMICAL ABSTRACTS, vol. 101, no. 13, 1984 Columbus, Ohio, US; abstract no. 110812c, SAAKYAN, A. M. ET AL.: "studies on the chlorination of organic compounds...." page 647; column 2; XP002218614 abstract & ARM. KHIM. ZH., vol. 37, no. 4, 1984, pages 261-265, ---	9
X	CHEMICAL ABSTRACTS, vol. 95, no. 1, 1981 Columbus, Ohio, US; abstract no. 6707u, AGARWAL, S. K. ET AL.: "Synthesis, characterization and screening of antibacterial activity of some new Mannich bases." page 635; column 1; XP002218615 abstract & J. INDIAN CHEM. SOC., vol. 57, no. 12, 1980, pages 1240-1241, ---	9
X	CHEMICAL ABSTRACTS, vol. 70, no. 9, 1969 Columbus, Ohio, US; abstract no. 37630b, TILAK, B. D. ET AL.: "Synthesis of nitrogen heterocyclics." page 331; column 1; XP002218616 abstract & INDIAN J. CHEM., vol. 6, no. 8, 1968, pages 422-427, ---	9

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INTERNATIONAL SEARCH REPORT

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C.(Continuation) DOCUMENTS CONSIDERED TO BE RELEVANT		
Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
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A	W. LEWIS ET AL.: "Ketonic Mannich bases....." JOURNAL OF THE AMERICAN PHARMACEUTICAL ASSOCIATION, vol. 47, no. 2, 1958, pages 77-81, XP001115095 page 80, column 2 -page 81, column 2 -----	9

INTERNATIONAL SEARCH REPORT

Information on patent family members

International Application No

PCT/EP 03/07411

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